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**The role of natural ingredients in anti-ageing of the skin**

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## **Introduction**

The skin ages for a number of reasons. It will naturally age with increasing loss of flexibility and ageing as collagen and elastin within the epidermis slowly cross-links and become less elastic. To a degree this is part of the genetic inheritance present within all of us, since do not seem to age at the same rate, nor share identical lifestyles.

It has been extensively proven that sunlight hastens the degradation of the skin by the bombardment of tissue with high energy photons present in UV-A and UV-B wavelengths of sunlight. This high energy has sufficient power to cleave molecules into free radicals, which are then available to react, modify and sometimes destroy healthy cellular chemistry.

Other external factors, such as the free radicals produced in vehicle exhaust gases, dirty industrial processes, and smoking can further speed the detriment of healthy skin.

## **Plants as skin protectants**

The role of plants in the protection of the skin may be come from a number of perspectives. Plants oils may be used to form a protective emollient layer that reduces transepidermal water loss and so increase the hydration of the stratum corneum. This not only forms a lubricious layer of fatty acids on the skin but also increases the “plumping” of the tissue, so contributing to a smoothing of the wrinkles.

The presence of reactive free radicals can be ‘mopped up’ by the use of antioxidants and free-radical scavengers.

The use of sunscreens will also reduce the potential of solar damage.

There is an increasing body of evidence to suggest that some plants can provide a prophylactic function.

The need to tan seems to be an irrepressible desire for many people and though one can tan safely without erythema and dangerous cellular damage, it is the feeling of most dermatologists that the ageing process is accelerated by tanning, regardless of the precautions that have been taken. The provision of cellular regeneration from plant sources is theoretically possible through the use of phytosterols and phytohormones and there are many other chemical entities within plants that can reduce erythema, reduce swelling and repair skin damage.

## **The chemistry of plant protectants**

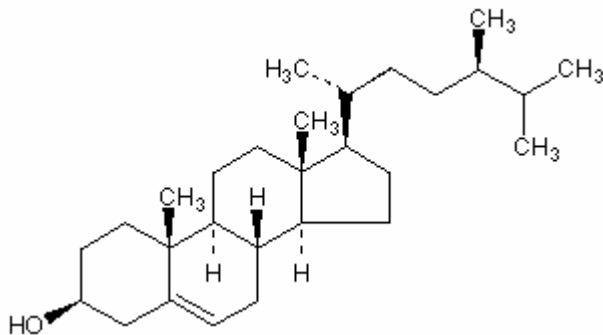
### ***Plant sterols***



## Plant sterols or phytosterols

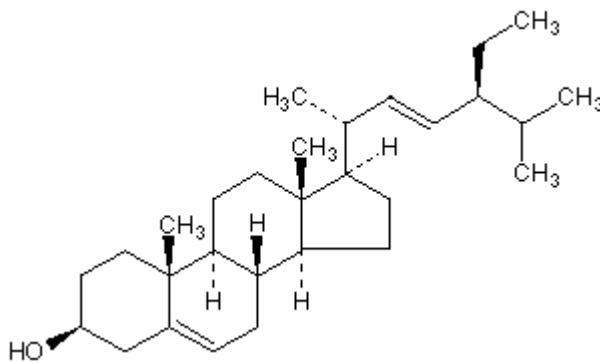
Another oil, Avocado (*Persea gratissima*), has also shown excellent effect on the skin,

particularly for the soothing, repairing and calming effect it exhibited. The analysis of the unsaponified components within this oil identified some phytosterols as the likely reason for this materials success on the skin as well as many other derivatives [Leung].



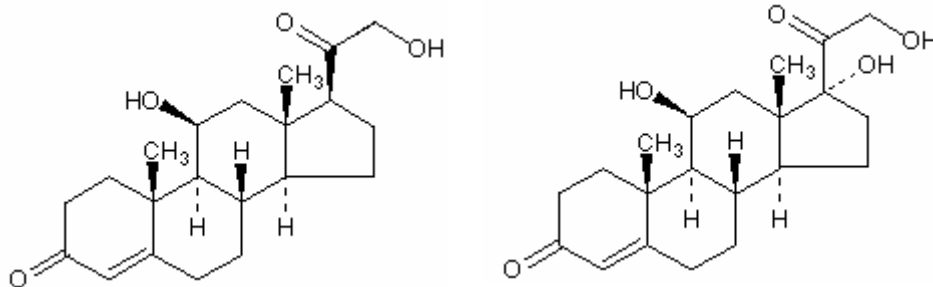
**Fig.4 Campesterol**

These included  $\beta$ -sitosterol, campesterol, stigmasterol, brassicasterol, delta5-avenasterol, tocopherols and other unidentified sterols [Crodarom].



These naturally occurring sterols bear a tremendous similarity to synthetic materials, those synthetic materials like corticosterone and hydrocortisone, traditionally used in allopathic medicine for their anti-inflammatory effects and particularly for skin eruptions, eczema and other pruritic and erythemic conditions.

**Fig.5 Stigmasterol**



**Fig.6 Corticosterone**

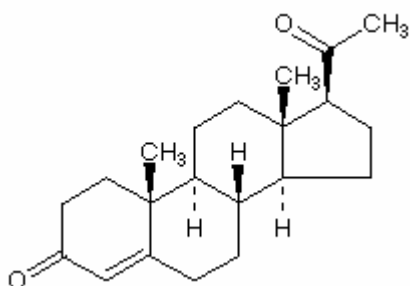
**Fig.7 Hydrocortisone**

Interestingly, the use of hydrocortisone or corticosteroids may ablate the symptoms of a distressing skin condition but not tackle the underlying reason for the skin reaction that underlies that condition. [Burden & Beck] Indeed, extended use of high levels of hydrocortisone can drastically alter the ratio between dermis, epidermis and stratum corneum and compromise the integrity of the skin to leave it with a thin topical fragility and visual transparency [Zachariae; Guin]. The answer to this apparent conundrum may lie within the chemistry, since though the structure bears an uncanny resemblance to the naturally occurring sterol materials, the exact configuration is

never found in nature. The skin may be “fooled” into reacting to the drug, because the synthetic steroid has a skeletal configuration or key that fits the lock.

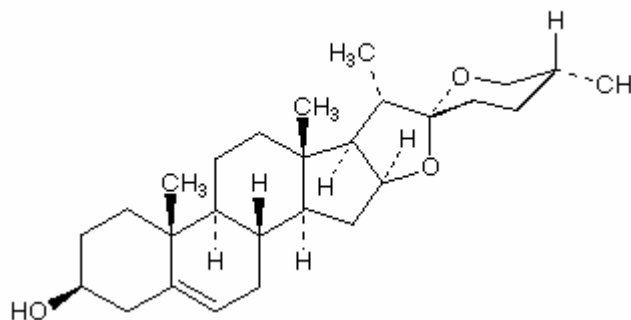
### ***Plant hormones***

In the old days (up until circa 1975) the cosmetic and toiletry industry were using oestrogen in products for mature skin until the pharmaceutical industry raised concerns that pure oestrogen was a pharmaceutical and it was subsequently banned. This material’s disappearance from the cosmetic ingredient inventory led to an outcry from the skin care manufacturers, but an even larger cry from the mature consumer who had discovered and proven for themselves the benefits of topical hormonal preparations.



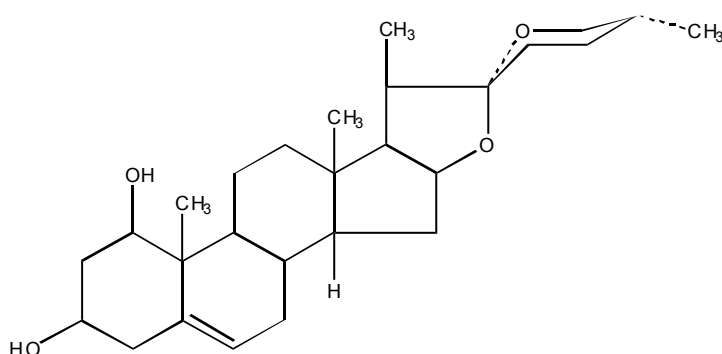
**Fig.8 Progesterone**

The pharmaceutical use that led to the ban was based on progesterone (one of the first female contraceptives), the raw material source was originally from Wild Yam (*Dioscorea villosa*) in the form of diosgenin.



**Fig.9 Diosgenin**

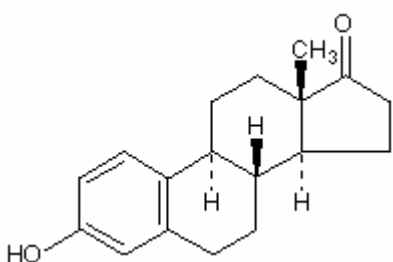
The diosgenin was converted to progesterone. Diosgenin is most certainly physiologically active and this can be seen by looking at molecule that is a close relative called ruscogenin (1-β-hydroxydiosgenin) and is found in Butcher’s Broom (*Ruscus aculeatus*) and used for the inflammatory conditions of oedema and associated erythema. It is also a useful material for the treatment of varicose veins and haemorrhoids.



**Fig. 10 Ruscogenin**

Eventually the stocks of Wild Yam fell to such low levels that industry was finding it hard to support the demand and so the much more widely available material Fenugreek (*Trigonella foenum-graecum*) was used as an alternative source of diosgenin. However the diosgenin by this time was being converted to another hormonal steroid, namely estrone.

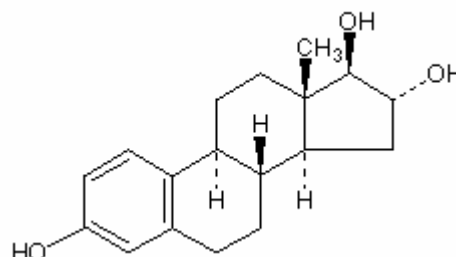
**Fig.11 Estrone**



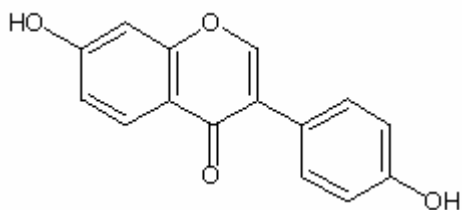
The use of plant materials as a source of these steroids faded as the products became produced synthetically. However, the relevance of plant steroids as a soothing and therapeutic source of active material is too irresistible to ignore.

A literature survey suggested that Ginseng (*Panax ginseng*) might be a source of estriol. This would be useful for the industry because there are very few papers published on the benefits of ginseng topically, because its main use is an alterative tonic taken internally.

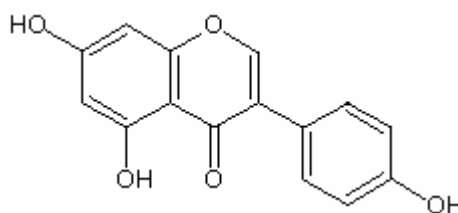
**Fig.12 Estriol**



Another plant material that was enjoying a great revival, until the issue of genetic modification spoilt the image, was soya (*Glycine max*). This material also contains  $\beta$ -sitosterol as illustrated above, but in addition contains daidzen and genistein, which are isoflavones (also anti-inflammatory and cell regenerating).

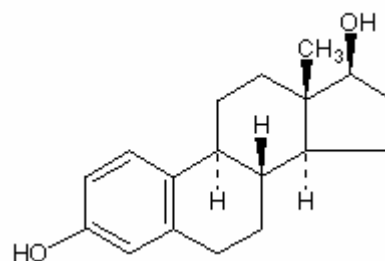


**Fig.13 Daidzein**



**Fig.14 Genistein**

The excitement of discovery grew immensely when we came to review a paper on *Pueraria mirifica* and discovered that this root also contained daidzein and genistein along with some other fascinating steroids and hormonal materials. In this case the steroidal material was estradiol and this offered an opportunity to offer another new steroidal derivative to our collection.



**Fig.15 Estradiol**

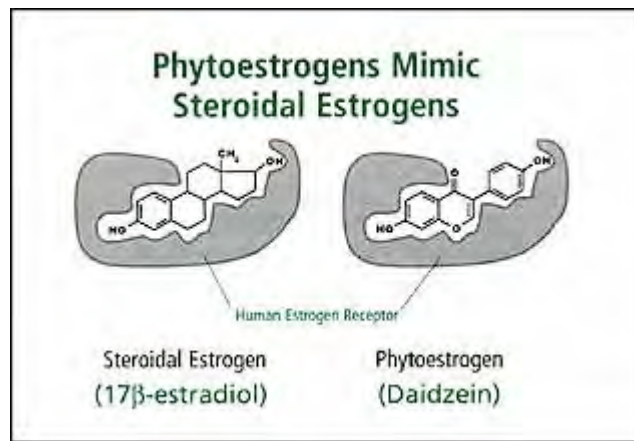
Proof of the steroidal nature of this Thai plant comes from its promotion as a bust developing preparation when taken internally. The supporting data is not conclusive, but shows some promise to its validity. Clearly this root would also have a topical application and benefit.

## Plant isoflavones

Isoflavones differ from flavones, because the phenyl group is attached to the 3-position, compared to flavones where the phenyl group is attached to the 2-position. The isoflavones occur naturally mainly within the Leguminosae (beans, soybeans, lentils, chick peas, etc).

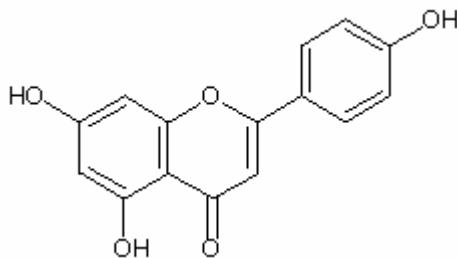
Well-known examples of isoflavones are genistein (4',5,7-trihydroxyisoflavone) and daidzein (4',7-dihydroxyisoflavone). Daidzein and genistein are phyto-estrogens. They are also named phenolic estrogens, to distinguish them from steroidal estrogens.

The geometrical structure of daidzein compares to 17- $\beta$ -estradiol and is therefore able to mimic its spatial structure [our thanks to Dr. Hans Brand for the use of his illustration]. Phyto-estrogens are much weaker in activity than steroidal estrogens, varying from 0.005 - 2%.



**Diag.1 molecular spatial representation**

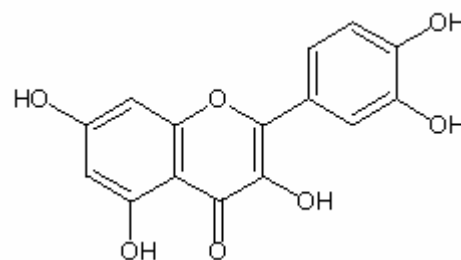
The estrogenic properties are by no means suitable to replace steroidal estrogens, but they do have significant interactions with the organism to enable to reduce the effects of ageing and include the improvement of the quality of the skin [Brand-Garnys].



**Fig.16 Apigenin**

The flavones, which are normally coloured [Dweck, 2002] also possess anti-inflammatory properties and are found in a variety of plants renowned for their soothing and healing properties.

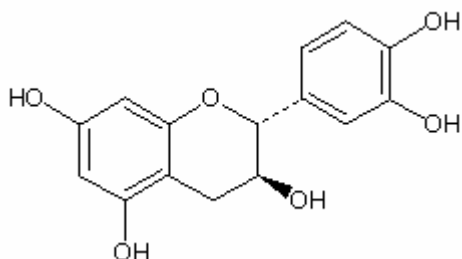
A closely related chemical group are the flavones, which include chemicals such as apigenin, luteolin and flavonols (which have an additional hydroxyl group in the 3-position) for example in quercetin.



**Fig.17 Quercetin**

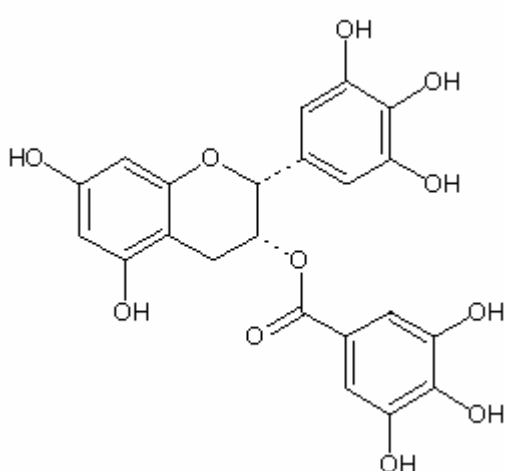
The iso-flavones have a phenyl group attached to the 3-position as opposed to the 2-position found in flavones. An example of an iso-flavone would be the genistein and daidzen already mentioned above.

These flavonoids have another important subgroup called the catechins, which have another remarkable property within plants and that is to be able to bind together in various configurations in order to form a powerful new complex group called procyanidins, which are free-radical scavengers and anti-oxidants.



**Fig. 18 Catechin**

Catechins lack the carbonyl group at the 4-position.



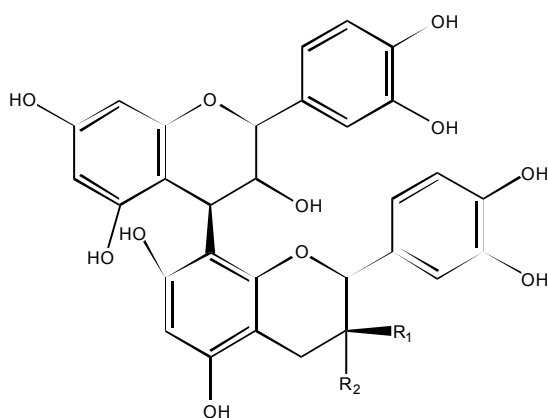
The formation of a powerful dimer called epigallocatechin-3-O-gallate occurs via the hydroxyl group and is found in materials like green tea (*Camelia sinensis*) and more recently in the stronger preparation white tea that is produced from the newly forming buds of that plant.

These materials (catechin and epicatechin) have also been extracted from the seeds of apples (*Pyrus malus*).

**Fig. 19 Epigallocatechin-3-O-gallate**

The wide distribution of these chemical moieties in nature suggests that the future will show a great opportunity for natural producers to extract and provide an even greater variety of these materials from ever more diverse sources.

### *Plant procyanidins*



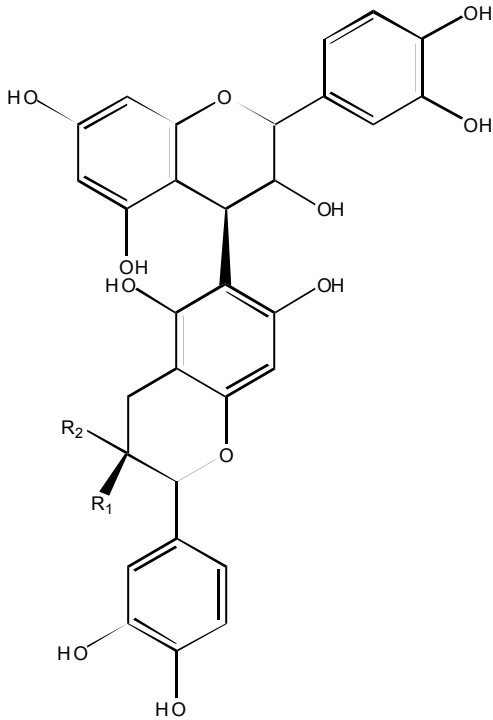
The classic source for the procyanidins has been the grape seed (*Vitis vinifera*), although a source was produced from marine pine and soya, that found its way into dietary supplements under the name pycnogenol (a name often used incorrectly for procyanidins) [Passwater and Kandaswami]

**Fig.20 Procyanidin B<sub>1</sub> and B<sub>2</sub>**

R<sub>1</sub> = OH; R<sub>2</sub> = H for Procyanidin B<sub>1</sub>  
R<sub>1</sub> = H; R<sub>2</sub> = OH for Procyanidin B<sub>2</sub>

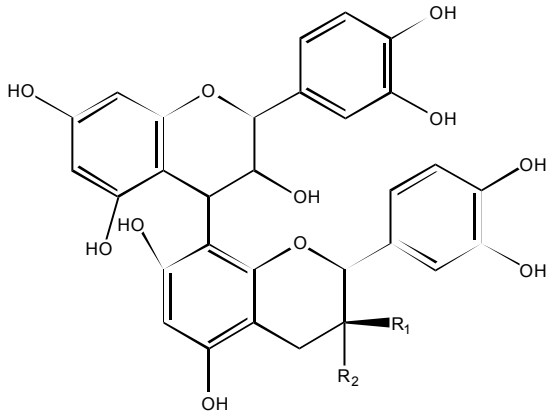
**Fig.21 Procyanidin B<sub>3</sub> and B<sub>4</sub>**

R<sub>1</sub> = OH; R<sub>2</sub> = H for Procyanidin B<sub>1</sub>  
R<sub>1</sub> = H; R<sub>2</sub> = OH for Procyanidin B<sub>2</sub>



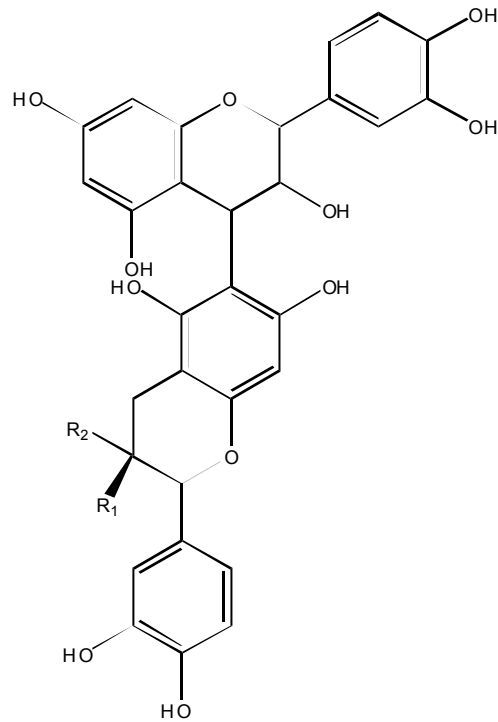
**Fig.22 Procyanidin B<sub>5</sub> and B<sub>7</sub>**

R<sub>1</sub> = OH; R<sub>2</sub> = H for Procyanidin B<sub>1</sub>  
R<sub>1</sub> = H; R<sub>2</sub> = OH for Procyanidin B<sub>2</sub>



**Fig.21 Procyanidin B<sub>6</sub> and B<sub>8</sub>**

R<sub>1</sub> = OH; R<sub>2</sub> = H for Procyanidin B<sub>1</sub>  
R<sub>1</sub> = H; R<sub>2</sub> = OH for Procyanidin B<sub>2</sub>



### ***Orchestral plants***

Up until now we have talked of single chemical species being responsible for the care and protection of the skin, in reality this is rarely the case in herbal medicine. Plants are not single chemical entities and do not give their maximum benefit when the single active molecule is extracted. The effect is not caused by a single instrument but more likely by an orchestra of different chemicals that work synergistically and in concert.

Two examples will be shown as a demonstration of this proposal.

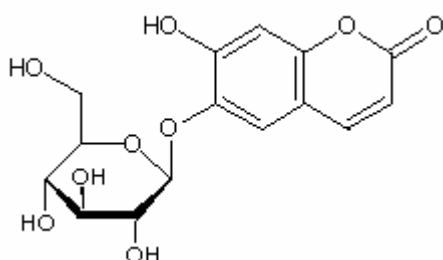
### ***Self Heal (*Prunella vulgaris*)***

Consider Self Heal (*Prunella vulgaris*) mentioned as long ago as around 1500AD by Theophrastus Bombastus von Hohenheim (better known to us as Paracelsus) in his *Doctrine of Signatures*.

It is a wound healing plant, used for swellings, pruritis and the treatment of cuts and abrasions. It has also been used by nursing mothers for sore and cracked nipples and is reputed to help stop the bleeding in wounds as an anti-haemorrhagic.

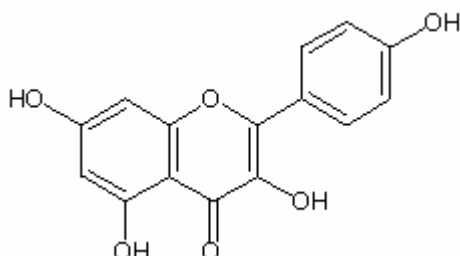
The chemical portfolio is quite remarkable and we find quercetin, esculin, delphinidin, rutin, aucubin, kaempferol and scopoletin.

One has to look at the root of the chemical name in order to gain a clue as to its original discovery and we find that esculin was originally called aesculin (and still is in some texts).



**Fig 23 Esculin**

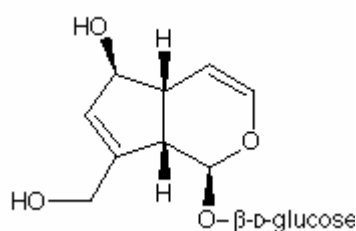
The root of the name aesculin is *Aesculus hippocastanum* or Horse Chestnut, which is well-known for its treatment of oedema and other inflammatory conditions [Fluck]. The esculin and related esculoside are attributed to with the properties of anti-oedema and anti-inflammatory effects [Weiss]. We can therefore make a guess that the esculin will exhibit similar properties in this plant.



We have already mentioned quercetin which is a flavonol (see Fig.13) and this is a well-respected soothing and healing agent, again with anti-inflammatory properties.

**Fig 24. Kaempferol**

Another flavonol present is kaempferol. This occurs widely in the plant world (often present with quercetin) and is found in a huge variety of plants that include Horse Chestnut (*Aesculus hippocastanum*), Arnica (*Arnica Montana*), Marigold (*Calendula officinalis*), Indian Pennywort (*Centella asiatica*) [Dweck, 1996] and Purple Coneflower (*Echinacea purpurea*) to name but a few. The plants have the same property in common - they are all renowned for their skin healing effect.

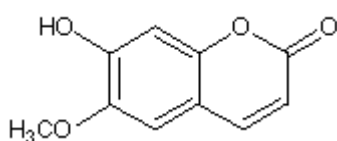


**Fig 25. Aucubin**

The root of this chemical would seem to come from *Aucuba japonica* – a Japanese variegated Laurel and chemically it is an iridoid glycoside. Aucubin is quite a difficult chemical to find data on. It is found in Eyebright (*Euphrasia officinalis*), Common Plantain (*Plantago officinalis*) and *Eucommia ulmoides*. The effects seem to be indicated as anti-viral and also to increase skin cell turnover, but these properties are far from being accurately defined.

The derivative of the chemical name scopoletin – an hydroxycoumarin - comes from the *Scopolia japonica* and the chemical is found in plants like Wormwood (*Artemisa annua*), Helichrysum picardii [Puerta], Horse Chestnut (*Aesculus hippocastanum*) [Komissarenko], Borage or Starflower (*Borago officinalis*) [Gudej] and Mallow (*Malva sylvestris*) [Tosi] where the plant is cited for its anti-inflammatory activity.

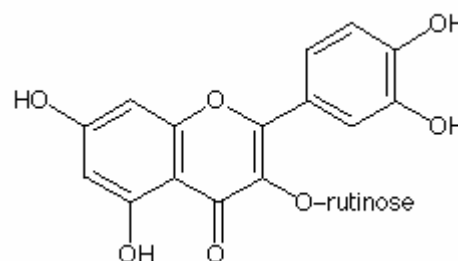
The exact function of scopoletin is hard to determine, some references lead towards it being a diuretic, while others lead to the reduction in PAF (Platelet Aggregation Factor). The likely action is anti-inflammatory from the studies on a variant of Cotton Lavender (*Santolina oblongifolia*) [Silvan] and also the antiinflammatory activity study of *Lomatia hirsuta* leaves (radal), a plant used in Chilean traditional medicine



for bronchial troubles and asthma, which was evaluated in guinea pigs by the carrageenan-induced paw oedema method [Erazo].

**Fig 26. Scopoletin**

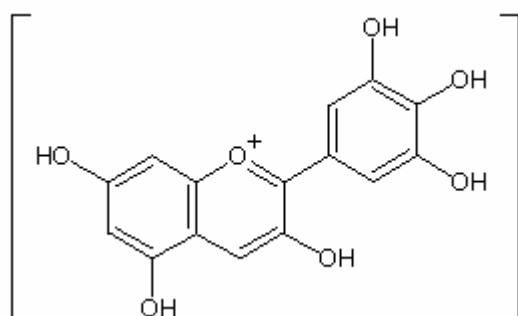
Rutin is found in the Graminae family and in products like Buckwheat (*Fagopyrum esculentum*) [Dabrowska-Zamojcin], *Anthriscus sylvestris* [Milovanovic] and *Ginkgo biloba* [Joyeux] where the effect appears to be antioxidant and protects cells against lipo-peroxidation. It is also found in plants like fennel (*Foeniculum vulgare*) and Aniseed (*Pimpinella anisum*).



**Fig 27. Rutin (Rutoside)**

Rutin or rutoside is shown as a capillary protectant [Merck]

The final material in this orchestra is a natural colour called delphinidin and closely related to flavones and flavanols.



**Fig 28 Delphinidin**

Its name is derived from the plant in which it was first found, namely delphinium and comes from the chemical family called anthocyanidins which coincidentally is another class of natural colourants [Dweck, 2002].

Another plant worthy of mention, that has recently come to attention is Tamanu oil or *Calophyllum inophyllum*. This has an extremely complex portfolio of chemistry that is hard to explain and decipher

## Tamanu (*Calophyllum inophyllum*)

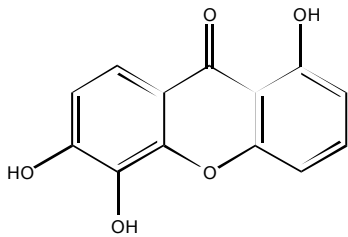
It is a member of the mangosteen family and the dried seeds produce a fixed oil. The full paper is published [Dweck, 2002;2]

Calophyllic acid and a lactone endowed with antibiotic properties are said to be at the origin of the oil's amazing cicatrising power. The dark-yellow oil extracted from the seeds contains a poisonous resin, which has a parsley-like odour. The resin is not dissimilar to myrrh and is alcohol-soluble.

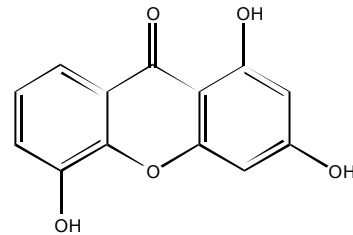
The bark contains tannin [Burkill] and exudes an oleoresin which contains benzoic acids [Jayaweera]. The oleoresin is officinal in the Mexican and Spanish Pharmacopoeias [Quisumbing].

The leaves of *Calophyllum inophyllum* contain friedelin and triterpenes of the friedelin group, namely canophyllal, canophyllol and canophyllic acid, [Govindachari, *et al*;1967] and from the heartwood xanthenes such as mesuaxanthone B and calophyllin B are obtained [Govindachari, *et al*;1968].

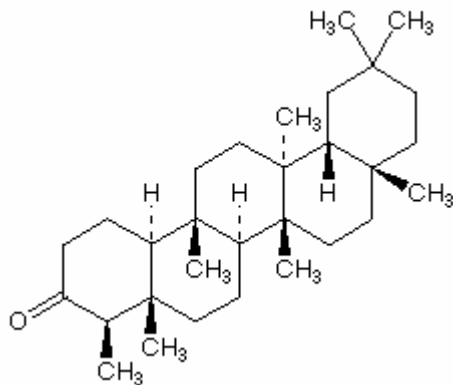
**Fig 29. Mesuaxanthone-B**



**Fig 30. Mesuaxanthone-A**



Interestingly, many of these substances i.e. canophyllal and friedelin are also found in *Syzygium* species.



**Fig 31. Friedelin**

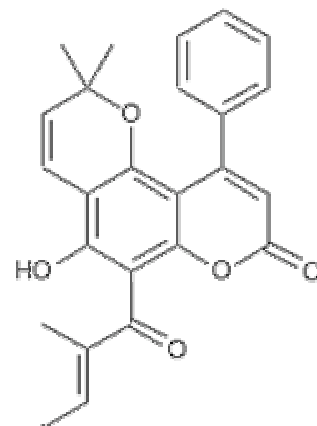
Friedelin has been identified. [Merck, 1996]

Calophyllolide, isolated from the nuts [Rastogi *et al*, 1998; Bhalla]. ED<sub>50</sub> was 140 mg/kg orally. Showed anti-inflammatory and anti-arthritic activity in formaldehyde-

induced arthritis and adjuvant arthritis in rats. LD<sub>50</sub> 2.5 g/kg p.o. It was devoid of ulcerogenic activity up to twice the ED<sub>50</sub> dose.

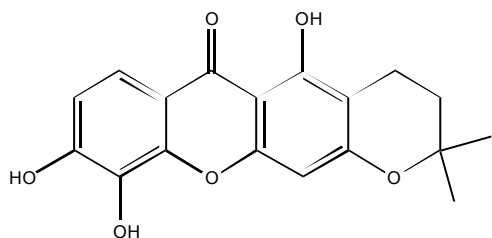
**Fig. 32 Calophyllolide**

De-hydrocycloguanindin, calophyllin-B, jacareubin and 6-deoxyjacareubin, produced varying degrees of CNS



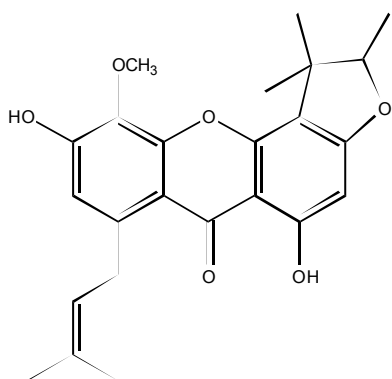
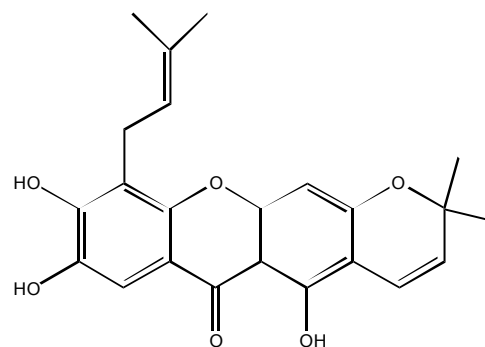
depression, characterised by ptosis, sedation, decreased spontaneous motor activity, loss of muscle tone, potentiation of phenobarbitone sleeping time and ether anaesthesia in mice and rats. All the xanthenes exhibited anti-inflammatory activity by both i.p. and oral routes in rats. Jacareubin and 6- deoxyjacareubin also showed anti-ulcer activity in rats [Gopalakrishnan *et al*].

**Fig. 33 Jacareubin**



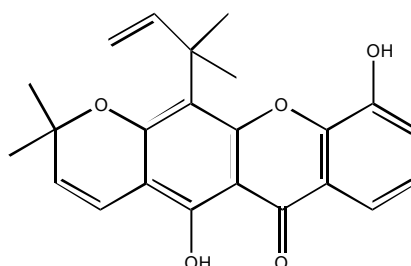
Together with inophyllide reduced oedema by 60.7% and 29.8% respectively (compared to hydrocortisone, which reduced inflammation by about 44%). The safety margin of calophyllolide is very similar to that of oxyphenbutazone (21.4 mg/kg and 25 mg/kg respectively).

Calophyllolide, isolated from the seeds [Oliver-Bever], reduced histamine inflammation and carrageenan-induced tissue swelling in rats.

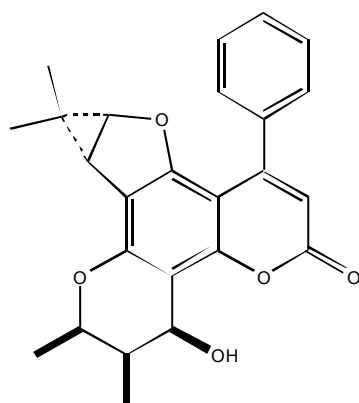


**Fig 35. Caloxanthone B**

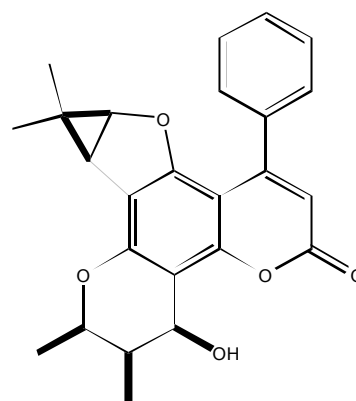
**Fig 34. Caloxanthone A**



**Fig 36. Caloxanthone C**



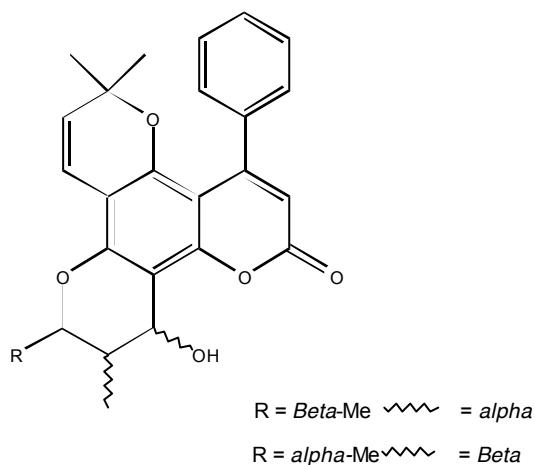
**Fig 37. Inophyllum G-1**



**Fig 38. Inophyllum G-2**

Inophyllums B and P inhibited HIV reverse transcriptase (IC<sub>50</sub> 38 and 130 nm respectively). [Rastogi *et al*, 1998]

Tamanu oil contains terpenic essences, benzoic and oxi-benzoic acids. Small amounts of vitamin F and phospho-amino-lipids come along with glycerides and saturated fatty acids.



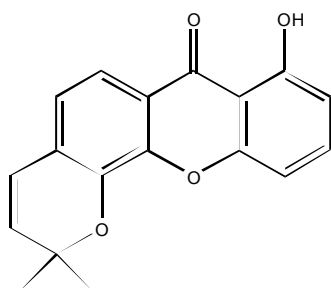
The plant contains 4-phenylcoumarins that have anti-tumour activity [Itoigawa]

**Fig 39. Inophyllum P**

The following active principles have been found in the oil:  
- calophyllolide (C<sub>25</sub>H<sub>22</sub>O<sub>5</sub>) the molecule of which contains a lactonic

and a methoxyl group.

- calophyllic acid (C<sub>25</sub>H<sub>24</sub>O<sub>6</sub>), which results from the saponification of the calophyllolide.



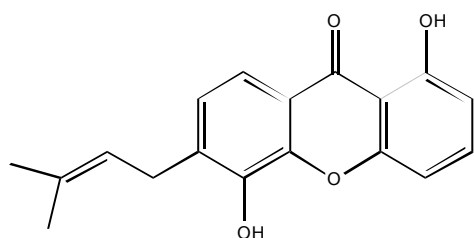
**Fig 40. Dehydrocycloguanandin**

These active principles are coumarin derivatives [Muller].  
Composition of the oil :

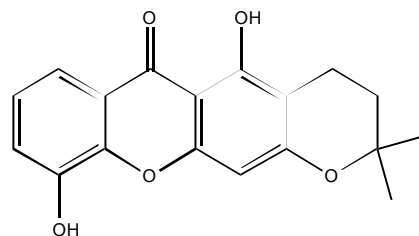
- \* Free fatty acids, glycerides, Sterols
- \* Terpenoids & steroids (canophyllal, canophyllol, calophyllic acid)

Coumarinic derivatives : Calophyllolids (natural neo-

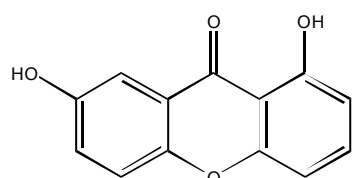
flavonoids with antibacterial, anti-inflammatory and anti-blood coagulation properties), Inophyllolids (natural neo-flavonoids with anti-viral properties), calophyllic acid (natural neo-flavonoid with anti-molluscidal and healing activities).



**Fig 41. Calaphyllin-B**

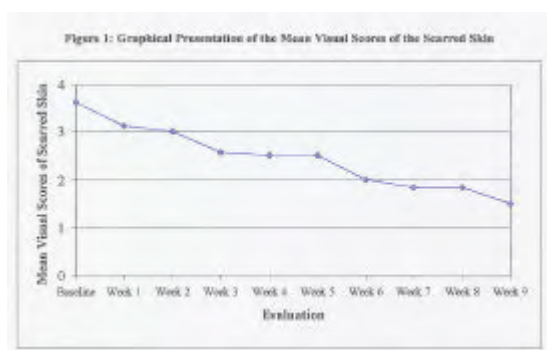
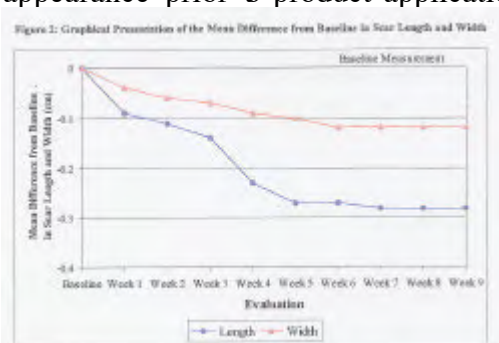


**Fig 42. 6-desoxyjacareubin**



**Fig 43. Euxanthone**

This study evaluated the ability of one (1) test product to improve the appearance of scars. Six (6) subjects with visually obvious, aged scars (1 year or more) were utilized for the study. The subjects were restricted from using any moisturizing products on the scarred area for a seven (7)-day pre-test period and throughout the nine (9) week test period. 0.5 mL aliquots of the product were applied to the scarred area twice a day for nine (9) consecutive weeks. Product applications were performed by the subjects and recorded on a product application tracking form provided to them. The subjects were evaluated prior to product application (baseline) and each week for nine (9) weeks at the testing facility. Visual ratings of scar appearance (color, roughness, and degree of difference from surrounding normal skin) and scar size measurements (length and width) were performed. Quantitative measurements of skin color for melanin (darkness) and hemoglobin (redness) were made on the scarred and adjacent normal skin areas using a Mexameter MX 18. Quantitative measurements of skin hydration were also performed on the same sites. Digital photographs of the scar were taken prior to product application (baseline) and again at the end of week nine (9). The subjects completed a self-evaluation questionnaire regarding their scar's appearance prior to product application



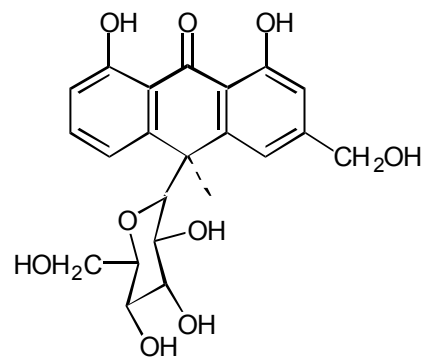
(baseline) and again at the end of week nine (9). The subjects also completed a product questionnaire that assessed their likes and dislikes of the product.

A significant improvement in the appearance of scars after six (6) weeks of Tamanu Oil use was observed visually. This improvement continued through to Week 9 of the study. The overall size of the scars consistently decreased throughout the study. The length of scars was reduced by an average of 0.28 cm, and the width by an average of 0.12 cm [Beausoleil].

### *Plants with prophylactic activity*

There is one plant that can most certainly protect the skin against radiation be it thermal [Fulton; Davis], solar [Strickland] or high energy radiowaves [Sato; Iena]. A review of the literature shows that the Aloe vera (*Aloe barbadensis*) can fulfil this promise [Reynolds & Dweck].

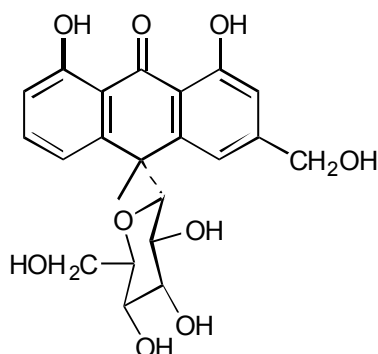
The chemistry is complex and the arguments still rage as to which component is responsible. The debate may never be resolved if researchers continue to look for just a single component responsible for the activity.



Barbaloin or Aloin A

**Fig 44. 10-(1',5'-anhydroglucosyl)-aloe-emodin-9-anthrone**

Isobarbaloin or Aloin B



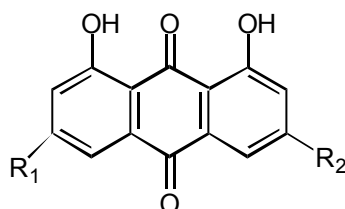
**Fig 45. 10-(1',5'-anhydroglucosyl)-aloe-emodin-9-anthrone**

These two chemicals are present as a mixture of:

10-C-β-L-glucosyldiastereo isomer of aloe-emodin anthrone and

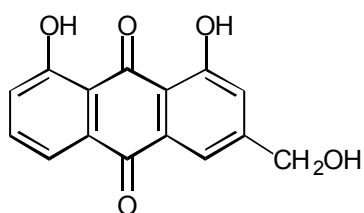
10-C-β-D-glucosyldiastereo isomer of aloe-emodin anthrone

They are present at 25-40% or 500mg/100g in the prepared aloe vera gel.



R <sub>1</sub>	R <sub>2</sub>	Chemical Name
H	CH <sub>3</sub>	CHRYSOPHANOL
H	CH <sub>2</sub> OH	ALOE-EMODOL
H	COOH	RHEINE
H	CH <sub>3</sub>	EMODOL

**Fig 46. Anthraquinone derivatives**



3-Hydroxymethyl anthraquinone

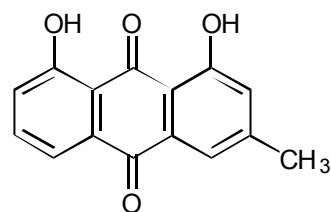
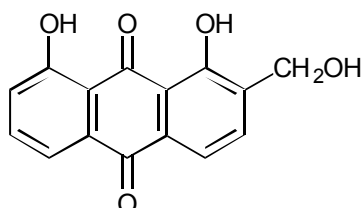
1,8-dihydroxy-3-hydroxymethyl-9,10-anthracenedione

Typically present at 2.05-2.2% in the aloe vera gel.

**Fig 47. Aloe-emodin or Aloe-emodol**

**Fig 48. Chrysophanol or Chrysarobin**

(Sometimes referred to Chrysophanic acid, which is a misnomer.)

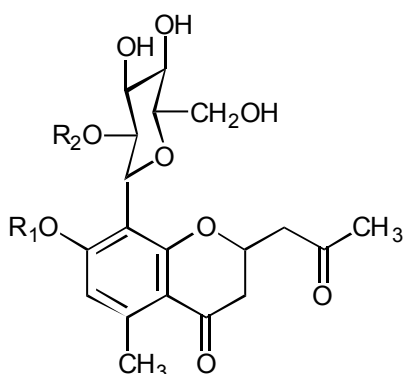


**Fig. 49 Aloetic acid**

Also mentioned is aloetic acid, which has a similar structure, but is not widely mentioned in the literature.

2-Hydroxymethylanthroquinone

## Chromone derivatives - Aloe resins



R <sub>1</sub>	R <sub>2</sub>	Chemical Name
H	<i>p</i> -coumaroyl	ALOERESIN A
H	H	ALOERESIN B
Glucosyl	<i>p</i> -coumaroyl	ALOERESIN C

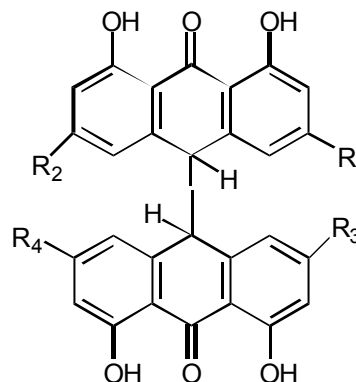
**Fig 50. Aloeresin A, Aloeresin B and Aloeresin C**  
 Aloeresin A is 2-*p*-coumaroyl aloeresin  
 Aloeresin B is 8-C-glucosylchromone aloeresin B  
 Aloeresin C is 7-O-β-D-glucoside of aloeresin

### Aloesone

The aglycone of aloeresins A, B and C. This is only present in trace amounts.

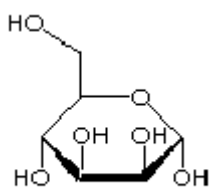
### Homodianthrones

**Fig 51. Sennidin A and B**



R <sub>1</sub>	R <sub>2</sub>	R <sub>3</sub>	R <sub>4</sub>	Chemical Name
COOH	H	COOH	H	SENNIDIN A, B
CH <sub>2</sub> OH	H	COOH	H	SENNIDIN C, D

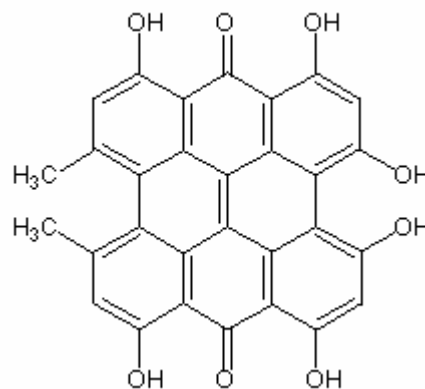
The above are the trace components, the major constituent is mannose-6-phosphate and it is this material which is attributed with the beneficial effects of the aloe vera. However, it is unlikely that this sugar derivative alone could give all the properties found for aloe in the scientific literature.



α-D-Mannose

The anthrones must play a part in the scenario a view perhaps shared by other authors [Joshi] who mention the chromones and anthracenes present possibly contributing to the action of the plant.

**Fig 52. Hypericin**



The clue may come from research into other medicinal plants, where it was proposed that the anthrone, anthraquinone, emodin, hypericin and pseudohypericin, isolated from St. John's Wort (*Hypericum perforatum*) may be involved in the anti-inflammatory effects [Bezakova]. The similarity between the structures of hypericin and sennidin (Fig.51) is too close to ignore.

## Conclusions

We have endeavoured to make a case for the protection of the skin using phytochemicals present in medicinal plants and have also speculated that the use of phytosterols and phytohormones would be beneficial to damaged and ageing skin. We examined those chemicals (from the published scientific literature) to the beneficial effects reported and were encouraged that the comparisons showed agreement.

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